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APPLICATION NO.	FILING DATE	FIRST NAMED INVENTOR	ATTORNEY DOCKET NO.	CONFIRMATION NO.
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09/833,711

04/13/2001

Luc Ouellet

10932-US

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CANADA

EXAMINER

TUROCY, DAVID P

ART UNIT

PAPER NUMBER

1762

MAIL DATE

DELIVERY MODE

06/20/2007

PAPER

**Please find below and/or attached an Office communication concerning this application or proceeding.**

The time period for reply, if any, is set in the attached communication.

**Office Action Summary**

Application No.

09/833,711

Applicant(s)

OUELLET ET AL.

Examiner

David Turocy

Art Unit

1762

-- The MAILING DATE of this communication appears on the cover sheet with the correspondence address --

**Period for Reply**

A SHORTENED STATUTORY PERIOD FOR REPLY IS SET TO EXPIRE 3 MONTH(S) OR THIRTY (30) DAYS, WHICHEVER IS LONGER, FROM THE MAILING DATE OF THIS COMMUNICATION.

- Extensions of time may be available under the provisions of 37 CFR 1.136(a). In no event, however, may a reply be timely filed after SIX (6) MONTHS from the mailing date of this communication.
- If NO period for reply is specified above, the maximum statutory period will apply and will expire SIX (6) MONTHS from the mailing date of this communication.
- Failure to reply within the set or extended period for reply will, by statute, cause the application to become ABANDONED (35 U.S.C. § 133). Any reply received by the Office later than three months after the mailing date of this communication, even if timely filed, may reduce any earned patent term adjustment. See 37 CFR 1.704(b).

**Status**

- 1) ☒ Responsive to communication(s) filed on 07 May 2007.
- 2a) ☒ This action is **FINAL**. 2b) ☐ This action is non-final.
- 3) ☐ Since this application is in condition for allowance except for formal matters, prosecution as to the merits is closed in accordance with the practice under *Ex parte Quayle*, 1935 C.D. 11, 453 O.G. 213.

**Disposition of Claims**

- 4) ☒ Claim(s) 1,3,6,8,14-18 and 21-23 is/are pending in the application.
- 4a) Of the above claim(s) \_\_\_\_\_ is/are withdrawn from consideration.
- 5) ☐ Claim(s) \_\_\_\_\_ is/are allowed.
- 6) ☒ Claim(s) 1,3,6,8,14-18 and 21-23 is/are rejected.
- 7) ☐ Claim(s) \_\_\_\_\_ is/are objected to.
- 8) ☐ Claim(s) \_\_\_\_\_ are subject to restriction and/or election requirement.

**Application Papers**

- 9) ☐ The specification is objected to by the Examiner.
- 10) ☐ The drawing(s) filed on \_\_\_\_\_ is/are: a) ☐ accepted or b) ☐ objected to by the Examiner.  
Applicant may not request that any objection to the drawing(s) be held in abeyance. See 37 CFR 1.85(a).  
Replacement drawing sheet(s) including the correction is required if the drawing(s) is objected to. See 37 CFR 1.121(d).
- 11) ☐ The oath or declaration is objected to by the Examiner. Note the attached Office Action or form PTO-152.

**Priority under 35 U.S.C. § 119**

- 12) ☐ Acknowledgment is made of a claim for foreign priority under 35 U.S.C. § 119(a)-(d) or (f).
- a) ☐ All b) ☐ Some \* c) ☐ None of:
1. ☐ Certified copies of the priority documents have been received.
  2. ☐ Certified copies of the priority documents have been received in Application No. \_\_\_\_\_.
  3. ☐ Copies of the certified copies of the priority documents have been received in this National Stage application from the International Bureau (PCT Rule 17.2(a)).

\* See the attached detailed Office action for a list of the certified copies not received.

**Attachment(s)**

- |  |   |
|--|---|
| 1) <input checked="" type="checkbox"/> Notice of References Cited (PTO-892)  | 4) <input type="checkbox"/> Interview Summary (PTO-413)<br>Paper No(s)/Mail Date. _____ |
| 2) <input type="checkbox"/> Notice of Draftsperson's Patent Drawing Review (PTO-948)                                   | 5) <input type="checkbox"/> Notice of Informal Patent Application (PTO-152)             |
| 3) <input type="checkbox"/> Information Disclosure Statement(s) (PTO-1449 or PTO/SB/08)<br>Paper No(s)/Mail Date _____ | 6) <input type="checkbox"/> Other: _____  |

## **DETAILED ACTION**

### ***Response to Amendment***

1. Applicants amendments, filed 5/7/2007, have been considered and reviewed by the examiner. The examiner notes the amendments to claim 1. Claims 1, 3, 6, 8, 14-18, 21-23 remain pending in the instant application.

### ***Response to Arguments***

2. Applicant's arguments filed 5/7/2007 have been fully considered but they are not persuasive.

The applicant has argued against the combination of the patents in the rejection, applying *KSR INT'L Co. v. Teleflex inc.*, arguing that the patents fail to note an existing a known problem at the time of the invention for which there was an obvious solution encompassed by the claims as required by the supreme court decision. However, the examiner notes that the while the Supreme Court decision may result in a new standard for measuring obviousness, such a new standard does not eliminate the prior standards that are in place. Therefore, it is the examiners position that the that a test of obviousness is not an express suggestion of the claimed invention in any or all references, but rather what the references taken collectively would suggest to those of ordinary skill in the art presumed to be familiar with them (*In re Rosselet*, 146 USPQ 183).

In response to applicant's argument that the examiner's conclusion of obviousness is based upon improper hindsight reasoning, it must be recognized that

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any judgment on obviousness is in a sense necessarily a reconstruction based upon hindsight reasoning. But so long as it takes into account only knowledge which was within the level of ordinary skill at the time the claimed invention was made, and does not include knowledge gleaned only from the applicant's disclosure, such a reconstruction is proper. See *In re McLaughlin*, 443 F.2d 1392, 170 USPQ 209 (CCPA 1971).

In response to applicant's argument that Ngo is nonanalogous art, it has been held that a prior art reference must either be in the field of applicant's endeavor or, if not, then be reasonably pertinent to the particular problem with which the applicant was concerned, in order to be relied upon as a basis for rejection of the claimed invention. See *In re Oetiker*, 977 F.2d 1443, 24 USPQ2d 1443 (Fed. Cir. 1992). In this case, Ojha et al. discloses depositing SiO<sub>2</sub> using PECVD process but fails to disclose the PECVD process and Ngo discloses known and suitable PECVD process conditions for deposition of silicon dioxide films. The selection of something based on its known suitability for its intended use has been held to support a *prima facie* case of obviousness. *Sinclair & Carroll Co. v. Interchemical Corp.*, 325 U.S. 327, 65 USPQ 297 (1945).

The applicant has argued against the Hsieh reference stating that the reference is concerned with the electrical properties of the PECVD deposited SiO<sub>2</sub> rather than optical properties. The applicant has argued that the reference "is in no way concerned with optical films, which really represents an entirely different art." However, clearly the applicant is erroneous in the interpretation of Hsieh. The examiner notes Hsieh

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discloses at figures 2, 3 and 4 discloses varying process parameters have a distinct effect on the physical properties of the film, including the refractive index and the absorbance. Each of the refractive index and the absorbance are known in the art to be optical properties of the depositing film, see as evidence such in Wikipedia explanation of both refractive index and absorbance. Such evidence is supplied as a general teaching to rebut the applicant's position that Hsieh is in no way concerned with the optical properties of deposited film.

The applicant has argued that the mere statement in Hsieh with regards to the connection between physical properties and the chamber pressure does not provide guidance to one of ordinary skill in the art. However, the examiner notes that Hsieh does not merely make a statement with regards to the connection. Hsieh discloses physical property of the deposited film is modified by the pressure, reactive gas ratio, rf pressure, total flow rate of the reactants, and the substrate temperature. Additionally, as shown above, the reference discloses such physical properties includes the optical property of the deposited film. As shown in figure 2, the optical properties of the film are optimized by maintaining, chamber pressure, power, and substrate temperature, and adjusting the flow ratio and/or flow rate into the chamber. Therefore the examiner maintains that Hsieh suggest to one of ordinary skill in the art that adjusting the pressure directly effects the optical properties of the deposited film.

The applicant has argued that the examiner has overlooked the claimed inventions, stating that the invention is directed to obtaining high quality optical films by fixing the flow rate at constant and varying the deposition pressure and comparing the

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FTIR spectra and based on the comparison selecting the deposition pressure.

However, Hsieh suggests selecting the optimum process parameter consists of maintaining all other process parameters constant during the process and varying a single process parameter, then comparing the results to determine the optimum for the single process parameter. Therefore, while the Hsieh fails to explicitly disclose maintaining a constant flow rate but varying the chamber pressure to determine the optimum film properties, such a modification would have been obvious to one of ordinary skill in the art because Hsieh discloses chamber pressure and total gas flow rate into the chamber both effect the physical properties of the deposited film (including optical properties such as absorbance and refractive index as evidenced by the Figures 2-4) and discloses adjusting a single process parameter, while maintaining the other parameters and then comparing the results to obtain the optimum deposited film.

### ***Claim Rejections - 35 USC § 103***

3. The following is a quotation of 35 U.S.C. 103(a) which forms the basis for all obviousness rejections set forth in this Office action:

(a) A patent may not be obtained though the invention is not identically disclosed or described as set forth in section 102 of this title, if the differences between the subject matter sought to be patented and the prior art are such that the subject matter as a whole would have been obvious at the time the invention was made to a person having ordinary skill in the art to which said subject matter pertains. Patentability shall not be negated by the manner in which the invention was made.

4. This application currently names joint inventors. In considering patentability of the claims under 35 U.S.C. 103(a), the examiner presumes that the subject matter of

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the various claims was commonly owned at the time any inventions covered therein were made absent any evidence to the contrary. Applicant is advised of the obligation under 37 CFR 1.56 to point out the inventor and invention dates of each claim that was not commonly owned at the time a later invention was made in order for the examiner to consider the applicability of 35 U.S.C. 103(c) and potential 35 U.S.C. 102(e), (f) or (g) prior art under 35 U.S.C. 103(a).

5. Claims 1, 3, 8, 14-18 are rejected under 35 U.S.C. 103(a) as being unpatentable over Ojha et al. (USPN 5,904,491) in view of Ngo et al. (USPN 6,127,261) and further in view of Hsieh et al. ("Characteristics of low-temperature and low-energy plasma-enhanced chemical vapor deposited SiO<sub>2</sub>", 1993).

Ojha et al. teaches a method of depositing an optical quality silica film on a substrate of a silica waveguide in manufacturing of a optical demultiplexer (Col.1, lines 3 – 8, Col.2, lines 43 – 58, Col. 5, lines 2019-20, and Col.6, lines 5 – 15), the method comprising forming an optical quality (i.e., silica) film on a substrate by PECVD in the presence of gases (Col.1, lines 64 – 67, Col.2, lines 1 – 10 and 43 – 57, Col.3, lines 10 – 11, 20 – 24, and 32 – 44, and Col.4, lines 22 – 29), and subjecting the as-deposited film to a low temperature treatment between 400° C to 1200° C, specifically at 800° C, to minimize the presence of contaminant compounds in the film (Col.1, lines 12 – 53 and 64 – 67, Col.2, lines 1 – 10 and 58 – 65, Col.3, lines 52 – 62, and Col.5, lines 20 – 36). While Ojha exemplified BPSG silica films, Ojha clearly discloses the cladding layer

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can be free of boron or phosphorus by alternatively doping with germania, titania, or arsenic (Column 3, lines 36-40).

Ojha et al. does not teach the specifics of the PECVD process, such as (1) a silane flow rate of 0.2 SLM, an N<sub>2</sub>O flow rate of 6.0 SLM, and a N<sub>2</sub> flow rate of 3.15 SLM, (2) a total deposition pressure of about 2.4 Torr, and (3) a deposition temperature of between 100 and 650° C, particularly 400° C. In general, Ojha et al. is silent as to the specifics of the PECVD process, except to say that the PECVD process may involve the use of silane and nitrous oxide as sources for silicon and oxygen, respectively, for the deposition of the waveguide material (Col.3, lines 19 – 24). Therefore, one of ordinary skill in the art would have been motivated to seek-out and utilize PECVD process parameters that are effective in depositing a silica film from gases such as silane and nitrous oxide, as desired by Ojha et al.

Ngo et al. teaches that the PECVD process recipe claimed by the applicant (i.e., a silane flow rate of 0.2 SLM, an N<sub>2</sub>O flow rate of 6.0 SLM, a N<sub>2</sub> flow rate of 3.15 SLM, a total deposition pressure of about 2.4 Torr, and a deposition temperature of between 100 and 650° C, particularly 400° C) is a well-known PECVD process recipe used to deposit silica films on a substrate (Col.4, lines 13 – 31, Claims 1 – 4). It would have been obvious to one of ordinary skill in the art to utilize the PECVD process parameters taught by Ngo et al. to deposit the silica film(s) of Ojha et al. because Ojha et al. generally desires to deposit a silica film by PECVD from silane and nitrous oxide reactants and Ngo et al. teaches a specific set of process parameters (e.g., gas flow



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rates, deposition pressure, deposition temperature, etc.) that are used to achieve just such a goal.

The combination of Ojha et al. and Ngo et al. does not explicitly teach that the annealing process minimizes the presence of Si-O-H-N compounds in the film. However, Ojha et al. teaches that the annealing process removes undesirable contaminants from the film in general (Col.1, lines 12 – 53 and 64 – 67, Col.2, lines 1 – 10 and 58 – 65, Col.3, lines 52 – 62, and Col.5, lines 20 – 36), and the combination of Ojha et al. and Ngo et al. teaches each and every process step and limitation of the applicant's claims, including the PECVD deposition temperature, the types of process gases, the deposition pressure, and the annealing temperature. As such, the annealing process of Ojha et al. would have inherently minimized the presence of Si-O-H-N compounds in the film, as claimed by the applicant. Please note that the mere observation of still another beneficial result (i.e., that annealing a PECVD silica film specifically reduces Si-O-H-N contaminants in the film, as opposed to contaminants in general) of an old process cannot form the basis of patentability (*Allen et al. v Coe*, 57 USPQ 136).

The combination of Ojha et al. and Ngo et al. does not explicitly teach the optimization process as claimed, however, Hsieh et al. teaches that, in the art of depositing silica films by PECVD from silane and N<sub>2</sub>O reactants (i.e., a process analogous to that of Ojha et al. and Ngo et al.), the more important variables that affect the electrical and physical properties of the deposited films include total flow rate of the reactants and chamber pressure (page 2639, col.2). Therefore Hsieh discloses the

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chamber pressure is a known result effective variable. The chamber pressure during deposition has a direct correlation on the physical properties of the deposited silica film by PECVD. Therefore it would have been obvious to one skill in the art at the time of the invention was made to determine the optimal value for the deposition pressure used in the process of Ojha et al. and Ngo et al. in view of Hsieh through routine experimentation, to impart the deposited film with the desired physical properties. Ojha et al. in view of Ngo et al. and Hsieh does not explicitly teach that the total deposition pressure is controlled to minimize the presence of Si-O-H-N compounds as evidenced by the FTIR characteristics. However, the total pressure taught by Ojha et al. in view of Ngo et al. (e.g., 2.0 to 2.4 Torr – see Ngo) is in the preferred range of pressures disclosed and claimed by the applicant (see, for example, applicant's Claim 1). Therefore, since the total pressure taught by Ojha et al. in view of Ngo et al. and Hsieh is in the range of pressures claimed by the applicant, the total pressure in Ojha et al. in view of Ngo et al. and Hsieh is inherently controlled "to minimize the presence of Si-O-H-N compounds". Please note that the mere observation of still another beneficial result (i.e., that a certain range of PECVD silica deposition pressures provides reduced contamination of the silica film) of an old process cannot form the basis of patentability (*Allen et al. v Coe*, 57 USPQ 136). Hsieh et al. also teaches using FTIR to analyze the composition and chemical bonds present in the deposited silica films, including any Si-H, Si-N, Si-O-H, N-H, etc. (page 2638, col.2; page 2639, col.1; page 640, column 1; page 2642, col.2). Therefore, it would have been obvious to one of ordinary skill in the art to analyze the silica films deposited by Ojha et al. and Ngo et al. with FTIR in order

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to reap the benefits of doing so, such as determining the composition and chemical bonds present in the films.

Hsieh suggests selecting the optimum process parameter consists of maintaining all other process parameters constant during the process and varying a single process parameter, then comparing the results to determine the optimum for the single process parameter. Therefore, while the Hsieh fails to explicitly disclose maintaining a constant flow rate but varying the chamber pressure to determine the optimum film properties, such a modification would have been obvious to one of ordinary skill in the art because Hsieh discloses chamber pressure and total gas flow rate into the chamber both effect the physical properties of the deposited film (including optical properties such as absorbance and refractive index as evidenced by the Figures 2-4) and discloses adjusting a single process parameter, while maintaining the other parameters and then comparing the results to obtain the optimum deposited film.

Ngo et al. also teaches a deposition temperature of about 400° C (**Claim 8**) (Column 4, lines 13-31); the applicant's claimed gases (i.e., SiH<sub>4</sub>, N<sub>2</sub>O, and N<sub>2</sub>) and optimized flow rates (**Claims 14 – 18**) (Column 4, lines 13-31);

6. Claim 6 is rejected under 35 U.S.C. 103(a) as being unpatentable over Ojha et al. (USPN 5,904,491) in view of Ngo et al. (USPN 6,127,261) and further in view of Hsieh et al. ("Characteristics of low-temperature and low-energy plasma-enhanced chemical vapor deposited SiO<sub>2</sub>", 1993) in further view of Chandross et al.

The combination of Ojha et al., Ngo et al., and Hsieh teaches all the limitations of **Claim 6** as set forth above, except for a method wherein the pressure is maintained by a vacuum pump having a controllable pumping speed, and the total gas pressure is maintained by controlling the pumping speed.

Specifically, the combination Ojha et al., Ngo et al., and Hsieh is silent as to how the appropriate chamber pressure is maintained. Chandross et al. teaches that it was known in the art of silicon oxide deposition at the time of the applicant's invention to maintain the desired pressure in a vacuum chamber by controlling the pumping speed of a vacuum pump (Col.5, lines 48 – 60). Therefore, it would have been obvious to one of ordinary skill in the art to maintain the desired pressure in the vacuum chamber during the PECVD process by controlling the pumping speed of a vacuum pump, as taught by Chandross et al., with the reasonable expectation of successfully and advantageously maintaining the desired pressure (e.g., 2.4 Torr) by utilizing a well-known, conventional means of doing so.

7. Claims 21 and 23 are rejected under 35 U.S.C. 103(a) as being unpatentable over Ojha et al. (USPN 5,904,491) in view of Ngo et al. (USPN 6,127,261).

Regarding **Claims 21 and 23** Ojha et al. teaches a method of depositing an optical quality silica film on a substrate of a silica waveguide in manufacturing of a optical demultiplexer (Col.1, lines 3 – 8, Col.2, lines 43 – 58, Col. 5, lines 2019-20, and Col.6, lines 5 – 15), the method comprising forming an optical quality (i.e., silica) film on a substrate by PECVD in the presence of gases (Col.1, lines 64 – 67, Col.2, lines 1 – 10

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and 43 – 57, Col.3, lines 10 – 11, 20 – 24, and 32 – 44, and Col.4, lines 22 – 29), and subjecting the as-deposited film to a low temperature treatment between 400° C to 1200° C, specifically at 800° C, to minimize the presence of contaminant compounds in the film (Col.1, lines 12 – 53 and 64 – 67, Col.2, lines 1 – 10 and 58 – 65, Col.3, lines 52 – 62, and Col.5, lines 20 – 36). While Ojha exemplified BPSG silica films, Ojha clearly discloses the cladding layer can be free of boron or phosphorus by alternatively doping with germania, titania, or arsenic (Column 3, lines 36-40).

Ojha et al. does not teach the specifics of the PECVD process, such as (1) a silane flow rate of 0.2 SLM, an N<sub>2</sub>O flow rate of 6.0 SLM, and a N<sub>2</sub> flow rate of 3.15 SLM, (2) a total deposition pressure of about 2.4 Torr, and (3) a deposition temperature of between 100 and 650° C, particularly 400° C. In general, Ojha et al. is silent as to the specifics of the PECVD process, except to say that the PECVD process may involve the use of silane and nitrous oxide as sources for silicon and oxygen, respectively, for the deposition of the waveguide material (Col.3, lines 19 – 24). Therefore, one of ordinary skill in the art would have been motivated to seek-out and utilize PECVD process parameters that are effective in depositing a silica film from gases such as silane and nitrous oxide, as desired by Ojha et al.

Ngo et al. teaches that the PECVD process recipe claimed by the applicant (i.e., a silane flow rate of 0.2 SLM, an N<sub>2</sub>O flow rate of 6.0 SLM, a N<sub>2</sub> flow rate of 3.15 SLM, a total deposition pressure of about 2.4 Torr, and a deposition temperature of between 100 and 650° C, particularly 400° C) is a well-known PECVD process recipe used to deposit silica films on a substrate (Col.4, lines 13 – 31, Claims 1 – 4). It would have

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been obvious to one of ordinary skill in the art to utilize the PECVD process parameters taught by Ngo et al. to deposit the silica film(s) of Ojha et al. because Ojha et al. generally desires to deposit a silica film by PECVD from silane and nitrous oxide reactants and Ngo et al. teaches a specific set of process parameters (e.g., gas flow rates, deposition pressure, deposition temperature, etc.) that are used to achieve just such a goal.

The combination of Ojha et al. and Ngo et al. does not explicitly teach that the annealing process minimizes the presence of Si-O-H-N compounds in the film. However, Ojha et al. teaches that the annealing process removes undesirable contaminants from the film in general (Col.1, lines 12 – 53 and 64 – 67, Col.2, lines 1 – 10 and 58 – 65, Col.3, lines 52 – 62, and Col.5, lines 20 – 36), and the combination of Ojha et al. and Ngo et al. teaches each and every process step and limitation of the applicant's claims, including the PECVD deposition temperature, the types of process gases, the deposition pressure, and the annealing temperature. As such, the annealing process of Ojha et al. would have inherently minimized the presence of Si-O-H-N compounds in the film, as claimed by the applicant. Please note that the mere observation of still another beneficial result (i.e., that annealing a PECVD silica film specifically reduces Si-O-H-N contaminants in the film, as opposed to contaminants in general) of an old process cannot form the basis of patentability (*Allen et al. v Coe*, 57 USPQ 136).

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8. Claim 22 is rejected under 35 U.S.C. 103(a) as being unpatentable over Ojha et al. (USPN 5,904,491) in view of Ngo et al. (USPN 6,127,261), in further view of Chandross et al.

The combination of Ojha et al. and Ngo et al. teaches all the limitations of **Claim 22** as set forth above, except for a method wherein the pressure is maintained by a vacuum pump having a controllable pumping speed, and the total gas pressure is maintained by controlling the pumping speed.

Specifically, the combination of Ojha et al. and Ngo et al. is silent as to how the appropriate chamber pressure is maintained. Chandross et al. teaches that it was known in the art of silicon oxide deposition at the time of the applicant's invention to maintain the desired pressure in a vacuum chamber by controlling the pumping speed of a vacuum pump (Col.5, lines 48 – 60). Therefore, it would have been obvious to one of ordinary skill in the art to maintain the desired pressure in the vacuum chamber during the PECVD process by controlling the pumping speed of a vacuum pump, as taught by Chandross et al., with the reasonable expectation of successfully and advantageously maintaining the desired pressure (e.g., 2.4 Torr) by utilizing a well-known, conventional means of doing so.

### ***Conclusion***

9. **THIS ACTION IS MADE FINAL.** Applicant is reminded of the extension of time policy as set forth in 37 CFR 1.136(a).

A shortened statutory period for reply to this final action is set to expire THREE MONTHS from the mailing date of this action. In the event a first reply is filed within TWO MONTHS of the mailing date of this final action and the advisory action is not mailed until after the end of the THREE-MONTH shortened statutory period, then the shortened statutory period will expire on the date the advisory action is mailed, and any extension fee pursuant to 37 CFR 1.136(a) will be calculated from the mailing date of the advisory action. In no event, however, will the statutory period for reply expire later than SIX MONTHS from the mailing date of this final action.

Any inquiry concerning this communication or earlier communications from the examiner should be directed to David Turocy whose telephone number is (571) 272-2940. The examiner can normally be reached on Monday-Friday 8:30-6:00, No 2nd Friday.

If attempts to reach the examiner by telephone are unsuccessful, the examiner's supervisor, Timothy Meeks can be reached on (571) 272-1423. The fax phone number for the organization where this application or proceeding is assigned is 571-273-8300.

Information regarding the status of an application may be obtained from the Patent Application Information Retrieval (PAIR) system. Status information for published applications may be obtained from either Private PAIR or Public PAIR. Status information for unpublished applications is available through Private PAIR only. For more information about the PAIR system, see <http://pair-direct.uspto.gov>. Should you have questions on access to the Private PAIR system, contact the Electronic Business Center (EBC) at 866-217-9197 (toll-free).




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**FRED J. PARKER**  
**PRIMARY EXAMINER**